

Variable	Mean	SD	Min	Max	Skewness	Kurtosis	Normality
Age	35.5	10.5	18	65	0.1	3.2	0.95
Gender	1.2	0.4	1	2	0.0	3.0	0.98
Marital Status	1.5	0.5	1	3	0.2	3.1	0.96
Education	12.5	2.5	8	16	0.3	3.3	0.94
Income	1500	500	500	3000	0.4	3.4	0.93
Occupation	1.8	0.6	1	3	0.1	3.1	0.97
Health Status	1.2	0.4	1	2	0.0	3.0	0.98
Stress Level	2.5	1.0	1	4	0.2	3.2	0.95
Life Satisfaction	3.5	1.5	1	5	0.3	3.3	0.94
Resilience	2.8	1.2	1	4	0.1	3.1	0.97
Optimism	3.2	1.4	1	5	0.2	3.2	0.96
Emotional Stability	2.0	0.8	1	3	0.1	3.1	0.97
Self-Esteem	3.0	1.0	1	4	0.2	3.2	0.95
Life Purpose	2.5	1.0	1	4	0.1	3.1	0.97
Gratitude	3.0	1.2	1	4	0.2	3.2	0.96
Forgiveness	2.8	1.1	1	4	0.1	3.1	0.97
Empathy	3.5	1.3	1	5	0.2	3.2	0.96
Compassion	3.2	1.4	1	5	0.1	3.1	0.97
Kindness	3.0	1.2	1	4	0.2	3.2	0.95
Patience	2.8	1.1	1	4	0.1	3.1	0.97
Humility	2.5	1.0	1	4	0.1	3.1	0.97
Generosity	3.0	1.2	1	4	0.2	3.2	0.95
Modesty	2.8	1.1	1	4	0.1	3.1	0.97
Self-Control	3.5	1.3	1	5	0.2	3.2	0.96
Discipline	3.2	1.4	1	5	0.1	3.1	0.97
Perseverance	3.0	1.2	1	4	0.2	3.2	0.95
Endurance	2.8	1.1	1	4	0.1	3.1	0.97
Stamina	3.0	1.2	1	4	0.2	3.2	0.95
Strength	2.8	1.1	1	4	0.1	3.1	0.97
Power	3.5	1.3	1	5	0.2	3.2	0.96
Influence	3.2	1.4	1	5	0.1	3.1	0.97
Authority	3.0	1.2	1	4	0.2	3.2	0.95
Leadership	2.8	1.1	1	4	0.1	3.1	0.97
Management	3.0	1.2	1	4	0.2	3.2	0.95
Organization	2.8	1.1	1	4	0.1	3.1	0.97
Coordination	3.5	1.3	1	5	0.2	3.2	0.96
Teamwork	3.2	1.4	1	5	0.1	3.1	0.97
Collaboration	3.0	1.2	1	4	0.2	3.2	0.95
Partnership	2.8	1.1	1	4	0.1	3.1	0.97
Relationship	3.0	1.2	1	4	0.2	3.2	0.95
Connection	2.8	1.1	1	4	0.1	3.1	0.97
Network	3.5	1.3	1	5	0.2	3.2	0.96
Community	3.2	1.4	1	5	0.1	3.1	0.97
Society	3.0	1.2	1	4	0.2	3.2	0.95
Culture	2.8	1.1	1	4	0.1	3.1	0.97
Tradition	3.0	1.2	1	4	0.2	3.2	0.95
Heritage	2.8	1.1	1	4	0.1	3.1	0.97
Legacy	3.5	1.3	1	5	0.2	3.2	0.96
Reputation	3.2	1.4	1	5	0.1	3.1	0.97
Image	3.0	1.2	1	4	0.2	3.2	0.95
Brand	2.8	1.1	1	4	0.1	3.1	0.97
Identity	3.0	1.2	1	4	0.2	3.2	0.95
Character	2.8	1.1	1	4	0.1	3.1	0.97
Personality	3.5	1.3	1				

2 A-X-M-Y-B

4 A and B are each members independently selected from the group consisting of
5 substituted and unsubstituted aryl and substituted and unsubstituted
6 heteroaryl;

$$\begin{array}{c} \text{a bond} \quad , \quad -\text{O}(\text{CH}_2)_m- \quad , \quad -\overset{\text{R}^1}{\text{N}}-(\text{CH}_2)_m- \quad , \quad -(\text{CH}_2)_n- \quad , \quad \begin{array}{c} \text{R}^2 \quad \text{R}^3 \\ \diagdown \quad \diagup \\ \text{C}=\text{C} \end{array} \quad , \\ \\ -\text{C}\equiv\text{C}- \quad , \quad \begin{array}{c} -\text{C}- \\ || \\ \text{W} \end{array} \quad , \quad \begin{array}{c} -\text{C}- \\ \diagdown \quad \diagup \\ \text{R}^2 \quad \text{R}^4 \end{array} \quad \text{and} \quad -\overset{\text{R}^1}{\text{N}}-\begin{array}{c} \text{C}- \\ || \\ \text{W} \end{array} \quad ; \end{array}$$

10 the subscript m is 0, 1 or 2;

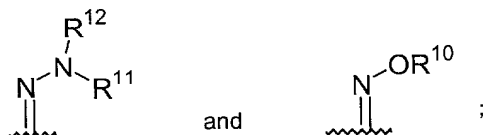
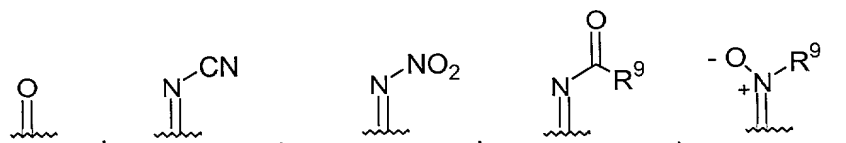
12 W is a member selected from the group consisting of O, N-OR⁵, N-NR¹R²,
13 N-NR¹C(O)R⁶ and N-OC(O)R⁶;

17 R⁴ is a member selected from the group consisting of H, OH, (C₁-C₆)alkyl,
18 (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino, di(C₁-C₆)alkylamino,
19 (C₁-C₆)acylamino, and (C₁-C₈)heteroalkyl; and

23 M is a divalent linking group selected from the group consisting of:

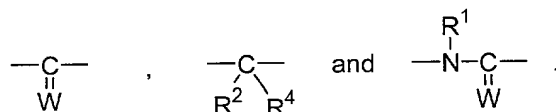
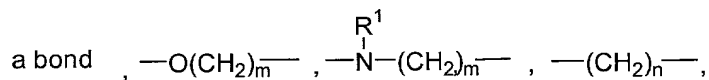
$$-\overset{\text{U}}{\underset{\text{C}}{\parallel}}-, \quad -\overset{\text{U}}{\underset{\text{C}}{\parallel}}-\overset{\text{R}^7}{\text{N}}-, \quad -\overset{\text{U}}{\underset{\text{C}}{\parallel}}-\underset{\text{R}^8}{\underset{\text{R}^9}{\text{C}}}-, \quad -\overset{\text{U}}{\underset{\text{C}}{\parallel}}-\overset{\text{O}}{\underset{\text{C}}{\parallel}}- \quad \text{and} \quad -\overset{\text{U}}{\underset{\text{C}}{\parallel}}-\overset{\text{C}}{\underset{\text{N}}{\parallel}}-\text{OR}^{10};$$

26 U is a member selected from the group consisting of:

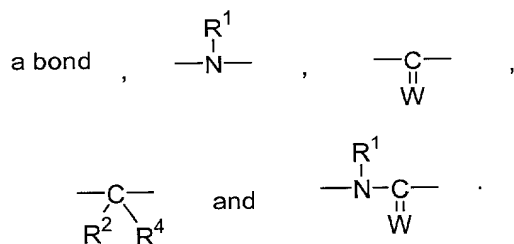


R^7 and R^8 are each independently members selected from the group consisting of H, OH, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino and di(C₁-C₆)alkylamino;
 R^9 is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl, aryl(C₁-C₆)alkyl, heteroaryl and heteroaryl(C₁-C₆)alkyl;
 R^{10} is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl and heteroaryl(C₁-C₆)alkyl; and
 R^{11} and R^{12} are members independently selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl, heteroaryl(C₁-C₆)alkyl, C(O)R¹⁴, C(O)OR¹⁴, C(O)-NR¹⁴R¹⁵, S(O)₂R¹³ and S(O)₂NR¹⁴R¹⁵;
 wherein
 R^{13} is a member selected from the group consisting of (C₁-C₆)alkyl, (C₁-C₆)heteroalkyl, phenyl and substituted phenyl; and
 R^{14} and R^{15} are each members independently selected from the group consisting of H, (C₁-C₆)alkyl and (C₁-C₆)heteroalkyl.

2. A compound of claim 1, wherein X and Y are independently selected from the group consisting of:



3. A compound of claim 1, wherein X and Y are each independently selected from the group consisting of:

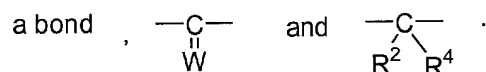


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4. A compound of claim 1, wherein X and Y are each independently selected from the group consisting of:



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1

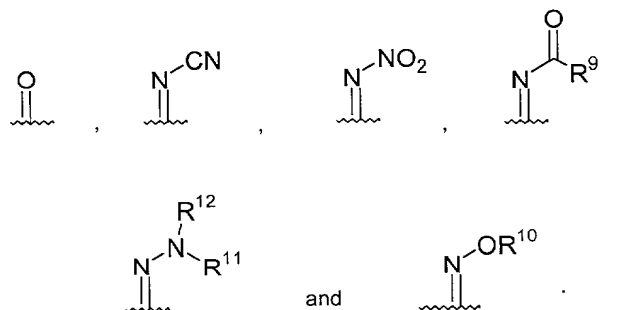
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5. A compound of claim 1, wherein M is $\text{—}\overset{\text{U}}{\parallel}\text{C}\text{—}\overset{\text{R}^7}{\text{N}}\text{—}$.

6. A compound of claim 1, wherein X and Y are each a bond, and M

is $\text{—}\overset{\text{U}}{\parallel}\text{C}\text{—}\overset{\text{R}^7}{\text{N}}\text{—}$, wherein U is selected from the group consisting of



3

1

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7. A compound of claim 6, wherein U is selected from the group consisting of

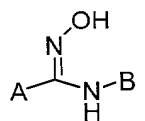


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8. A compound of claim 1, said compound having the formula:

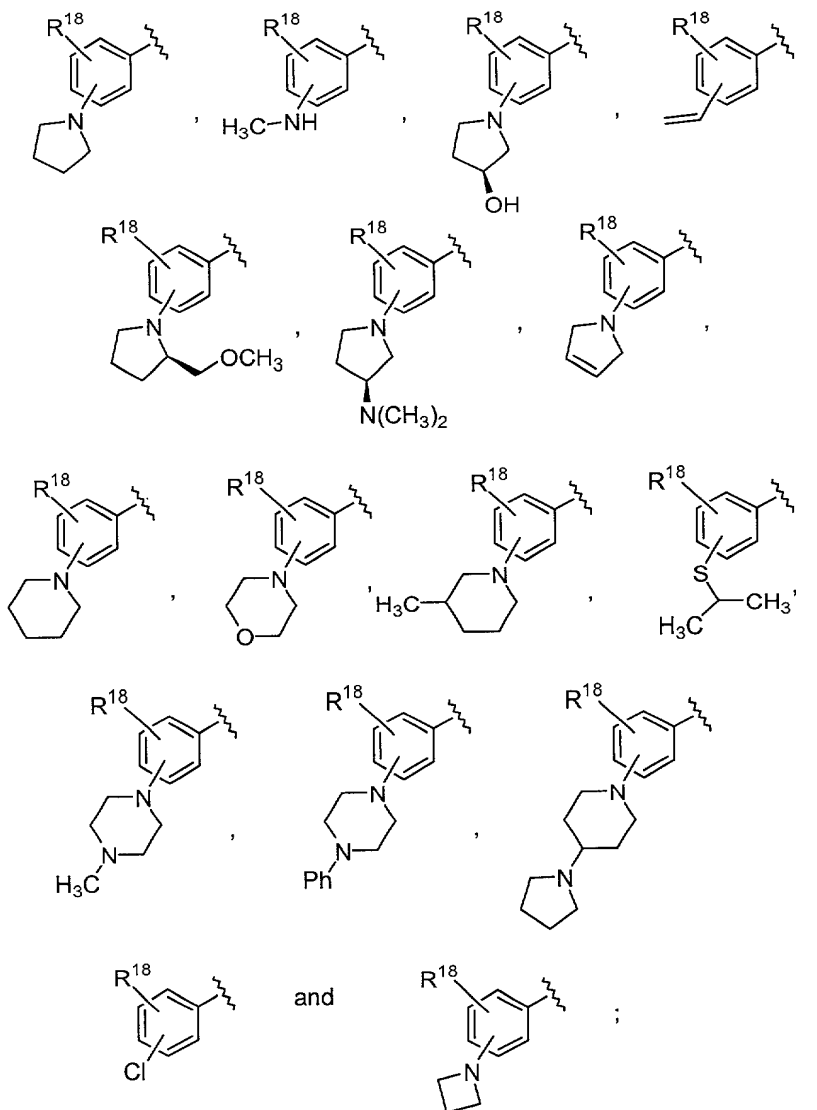


1 9. A compound of claim 8, wherein A is a phenyl group substituted
2 with from one to three substituents selected from the group consisting of (C₁-C₄)alkyl,
3 (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, nitro, phenyl, naphthyl,
4 pyrrolyl, pyrazolyl and -NR¹⁶R¹⁷ wherein R¹⁶ and R¹⁷ are independently selected from
5 the group consisting of hydrogen, (C₁-C₈)alkyl and (C₁-C₈)heteroalkyl or are combined
6 with the nitrogen atom to which each is attached to form a four-, five-, six- or seven-
7 membered ring optionally having additional heteroatoms as ring members and optionally
8 having additional substituents selected from the group consisting of (C₁-C₈)alkyl, (C₁-
9 C₈)heteroalkyl and phenyl.

1 10. A compound of claim 8, wherein B is a phenyl group substituted
2 with from one to three substituents selected from the group consisting of (C₁-C₄)alkyl,
3 (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, phenyl
4 and phenoxy.

1 11. A compound of claim 8, wherein A is a phenyl group substituted
2 with from one to three substituents selected from the group consisting of (C₁-C₄)alkyl,
3 (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen and -NR¹⁶R¹⁷ wherein R¹⁶
4 and R¹⁷ are independently selected from the group consisting of hydrogen, (C₁-C₈)alkyl
5 and (C₁-C₈)heteroalkyl or are combined with the nitrogen atom to which each is attached
6 to form a four-, five-, six- or seven-membered ring optionally having additional
7 heteroatoms as ring members and optionally having additional substituents selected from
8 the group consisting of (C₁-C₈)alkyl, (C₁-C₈)heteroalkyl and phenyl, and B is a phenyl
9 group substituted with from one to three substituents selected from the group consisting
10 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy,
11 halogen, phenyl and phenoxy.

1 12. A compound of claim 8, wherein A is selected from the group
2 consisting of substituted or unsubstituted thienyl, substituted or unsubstituted furanyl,
3 substituted or unsubstituted indolyl, substituted or unsubstituted benzothienyl, substituted
4 or unsubstituted benzothienyl, and radicals of the formulae:



5
6 wherein R¹⁸ is a member selected from the group consisting of (C₁-
7 C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy and
8 halogen.

1 **13.** A compound of claim 8, wherein A is selected from the group
2 consisting of substituted or unsubstituted benzofuranyl, substituted or unsubstituted
3 benzothienyl, substituted or unsubstituted indolyl, substituted or unsubstituted
4 benzimidazolyl, substituted or unsubstituted benzthiazolyl and substituted or
5 unsubstituted benzoxazolyl.

1 **14.** A method of reducing bacterial growth on a surface, said method
2 comprising contacting said surface with a compound of claim 1.

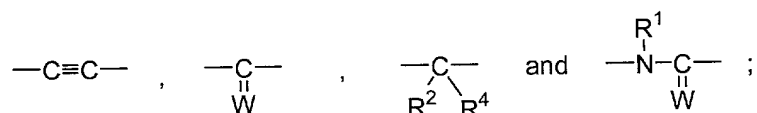
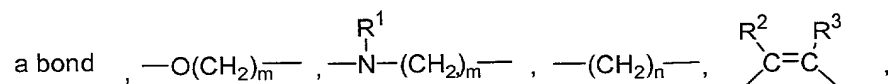
1 15. A method of treating a bacterial infection comprising contacting a
2 subject in need of such treatment with an effective amount of a compound having the
3 formula:



4
5 or a pharmaceutically acceptable salt thereof, wherein

6 A and B are each members independently selected from the group consisting of
7 substituted and unsubstituted aryl and substituted and unsubstituted
8 heteroaryl;

9 X and Y are each members independently selected from the group consisting of:



10

11 with the proviso that at least one of X or Y is a bond, and wherein

12 the subscript m is 0, 1 or 2;

13 the subscript n is 1 or 2;

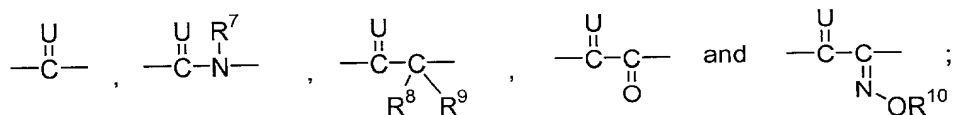
14 W is a member selected from the group consisting of O, N-OR⁵, N-NR¹R²,
15 N-NR¹C(O)R⁶ and N-OC(O)R⁶;

16 R¹, R², R³ and R⁵ are each members independently selected from the group
17 consisting of H, (C₁-C₆)alkyl, aryl, aryl(C₁-C₆)alkyl, heteroaryl and
18 heteroaryl(C₁-C₆)alkyl;

19 R⁴ is a member selected from the group consisting of H, OH, (C₁-C₆)alkyl,
20 (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino, di(C₁-C₆)alkylamino,
21 (C₁-C₆)acylamino, and (C₁-C₈)heteroalkyl; and

22 R⁶ is a member selected from the group consisting of H, (C₁-C₆)alkyl, (C₁-
23 C₆)alkoxy, amino, (C₁-C₆)alkylamino, di(C₁-C₆)alkylamino and
24 (C₁-C₈)heteroalkyl; and

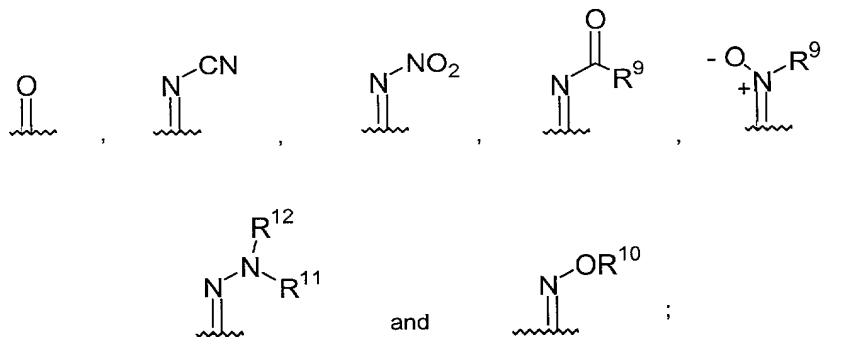
25 M is a divalent linking group selected from the group consisting of:



26

27 wherein

28 U is a member selected from the group consisting of:



R^7 and R^8 are each members independently selected from the group

consisting of H, OH, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino and di(C₁-C₆)alkylamino;

R^9 is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl, aryl(C₁-C₆)alkyl, heteroaryl and heteroaryl(C₁-C₆)alkyl;

R^{10} is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl and heteroaryl(C₁-C₆)alkyl; and

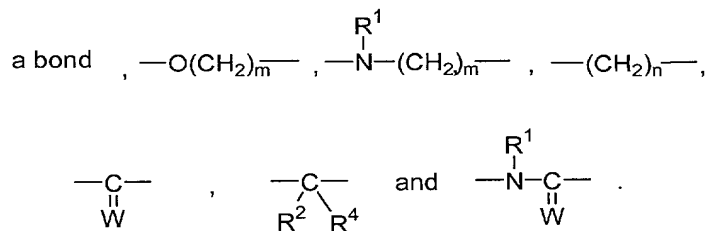
R^{11} and R^{12} are members independently selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl, heteroaryl(C₁-C₆)alkyl, C(O)R¹⁴, C(O)OR¹⁴, C(O)-NR¹⁴R¹⁵, S(O)₂R¹³ and S(O)₂NR¹⁴R¹⁵;

wherein

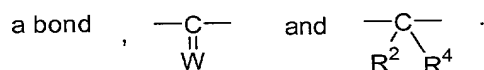
R^{13} is a member selected from the group consisting of (C₁-C₆)alkyl, (C₁-C₆)heteroalkyl, phenyl and substituted phenyl; and

R^{14} and R^{15} are each members independently selected from the group consisting of H, (C₁-C₆)alkyl and (C₁-C₆)heteroalkyl.

16. A method in accordance with claim 15, wherein X and Y are independently selected from the group consisting of:



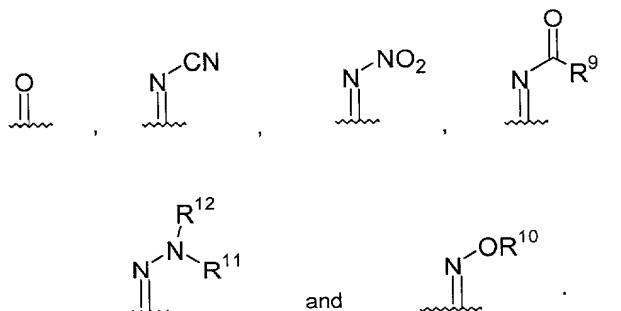
17. A method in accordance with claim 15, wherein X and Y are each independently selected from the group consisting of:



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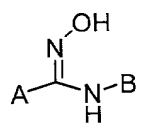
1 **18.** A method in accordance with claim **15**, wherein X and Y are each a

2 bond, and M is $\begin{array}{c} \text{U} \quad \text{R}^7 \\ || \quad | \\ \text{---C---N---} \end{array}$, wherein U is selected from the group consisting of



3

1 **19.** A method in accordance with claim **15**, said compound having the
2 formula:



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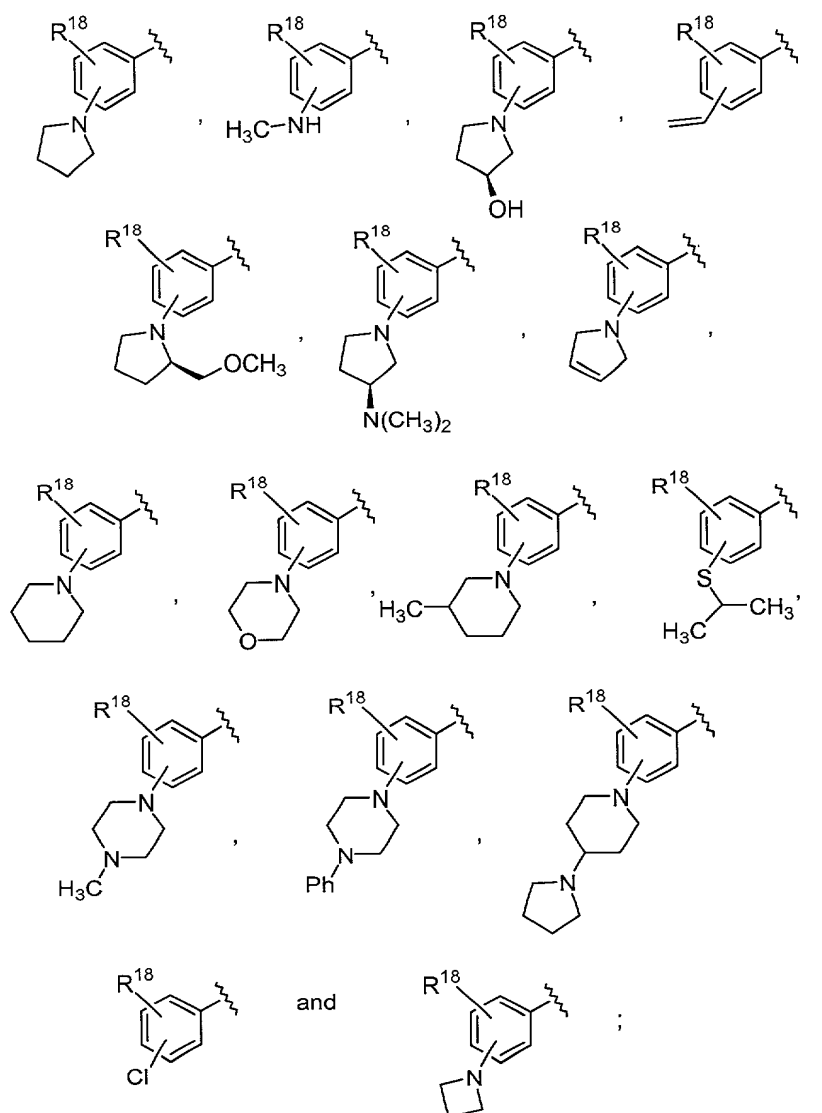
1 **20.** A method in accordance with claim **19**, wherein A is a phenyl
2 group substituted with from one to three substituents selected from the group consisting
3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, nitro,
4 phenyl, naphthyl, pyrrolyl, pyrazolyl and -NR¹⁶R¹⁷ wherein R¹⁶ and R¹⁷ are
5 independently selected from the group consisting of hydrogen, (C₁-C₈)alkyl and (C₁-
6 C₈)heteroalkyl or are combined with the nitrogen atom to which each is attached to form
7 a four-, five-, six- or seven-membered ring optionally having additional heteroatoms as
8 ring members and optionally having additional substituents selected from the group
9 consisting of (C₁-C₈)alkyl, (C₁-C₈)heteroalkyl and phenyl.

1 **21.** A method in accordance with claim **19**, wherein B is a phenyl
2 group substituted with from one to three substituents selected from the group consisting
3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy,
4 halogen, phenyl and phenoxy.

1 **22.** A method in accordance with claim **19**, wherein A is a phenyl
2 group substituted with from one to three substituents selected from the group consisting

3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen and –
4 NR¹⁶R¹⁷ wherein R¹⁶ and R¹⁷ are independently selected from the group consisting of
5 hydrogen, (C₁-C₈)alkyl and (C₁-C₈)heteroalkyl or are combined with the nitrogen atom to
6 which each is attached to form a four-, five-, six- or seven-membered ring optionally
7 having additional heteroatoms as ring members and optionally having additional
8 substituents selected from the group consisting of (C₁-C₈)alkyl, (C₁-C₈)heteroalkyl and
9 phenyl, and B is a phenyl group substituted with from one to three substituents selected
10 from the group consisting of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-
11 C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, phenyl and phenoxy.

1 **23.** A method in accordance with claim **19**, wherein A is selected from
2 the group consisting of substituted or unsubstituted thienyl, substituted or unsubstituted
3 furanyl, substituted or unsubstituted indolyl, substituted or unsubstituted benzothienyl,
4 substituted or unsubstituted benzothienyl, and radicals of the formulae:



5
6 wherein R^{18} is a member selected from the group consisting of (C₁-
7 C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy and
8 halogen.

1 **24.** A method in accordance with claim **23**, wherein A is selected from
2 the group consisting of substituted or unsubstituted benzofuranyl, substituted or
3 unsubstituted benzothieryl, substituted or unsubstituted indolyl, substituted or
4 unsubstituted benzimidazolyl, substituted or unsubstituted benzthiazolyl and substituted
5 or unsubstituted benzoxazolyl.

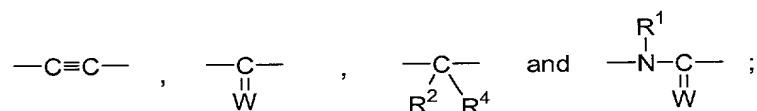
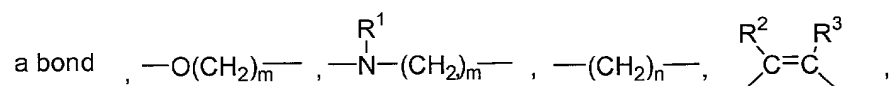
1 **25.** A composition comprising a pharmaceutically acceptable excipient
2 in admixture with a compound having the formula:



or a pharmaceutically acceptable salt thereof, wherein

A and B are each members independently selected from the group consisting of substituted and unsubstituted aryl and substituted and unsubstituted heteroaryl;

X and Y are each members independently selected from the group consisting of:



with the proviso that at least one of X or Y is a bond, and wherein

the subscript m is 0, 1 or 2;

the subscript n is 1 or 2;

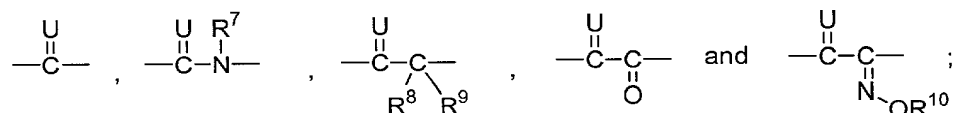
W is a member selected from the group consisting of O, N-OR⁵, N-NR¹R², N-NR¹C(O)R⁶ and N-OC(O)R⁶;

R¹, R², R³ and R⁵ are each members independently selected from the group consisting of H, (C₁-C₆)alkyl, aryl, aryl(C₁-C₆)alkyl, heteroaryl and heteroaryl(C₁-C₆)alkyl;

R⁴ is a member selected from the group consisting of H, OH, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, (C₁-C₆)acylamino, and (C₁-C₈)heteroalkyl; and

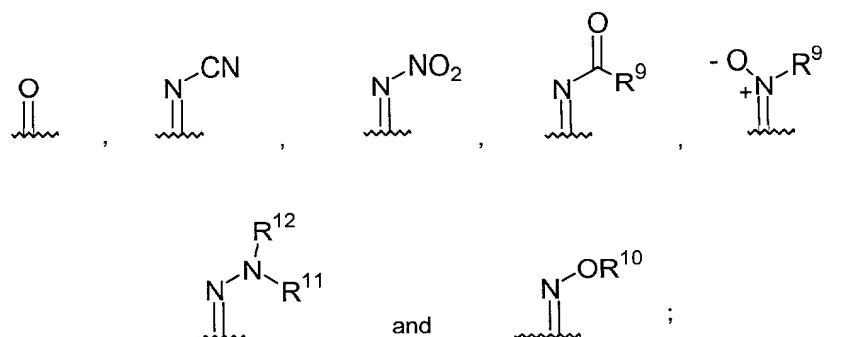
R⁶ is a member selected from the group consisting of H, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino, di(C₁-C₆)alkylamino and (C₁-C₈)heteroalkyl; and

M is a divalent linking group selected from the group consisting of:



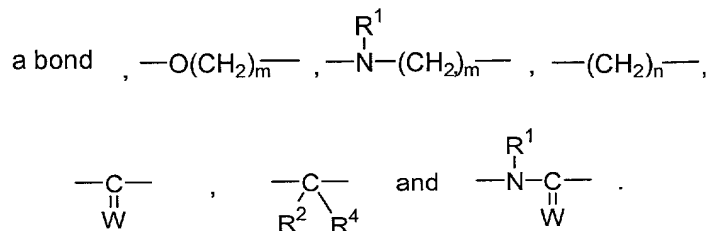
wherein

U is a member selected from the group consisting of:

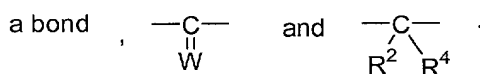


R^7 and R^8 are each members independently selected from the group consisting of H, OH, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, amino, (C₁-C₆)alkylamino and di(C₁-C₆)alkylamino;
 R^9 is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl, aryl(C₁-C₆)alkyl, heteroaryl and heteroaryl(C₁-C₆)alkyl;
 R^{10} is a member selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl and heteroaryl(C₁-C₆)alkyl; and
 R^{11} and R^{12} are members independently selected from the group consisting of H, (C₁-C₆)alkyl, aryl(C₁-C₆)alkyl, heteroaryl(C₁-C₆)alkyl, C(O) R^{14} , C(O)OR¹⁴, C(O)-NR¹⁴R¹⁵, S(O)₂R¹³ and S(O)₂NR¹⁴R¹⁵;
 wherein
 R^{13} is a member selected from the group consisting of (C₁-C₆)alkyl, (C₁-C₆)heteroalkyl, phenyl and substituted phenyl; and
 R^{14} and R^{15} are each members independently selected from the group consisting of H, (C₁-C₆)alkyl and (C₁-C₆)heteroalkyl.

26. A composition in accordance with claim **25**, wherein X and Y are independently selected from the group consisting of:



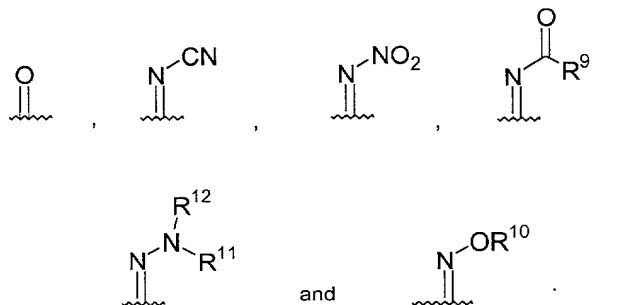
27. A composition in accordance with claim **25**, wherein X and Y are each independently selected from the group consisting of:



3

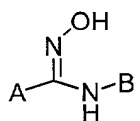
1 **28.** A composition in accordance with claim **25**, wherein X and Y are

2 each a bond, and M is $\begin{array}{c} \text{U} \quad \text{R}^7 \\ || \quad | \\ \text{---C---N---} \end{array}$, wherein U is selected from the group consisting of



3

1 **29.** A composition in accordance with claim **25**, said compound having
2 the formula:



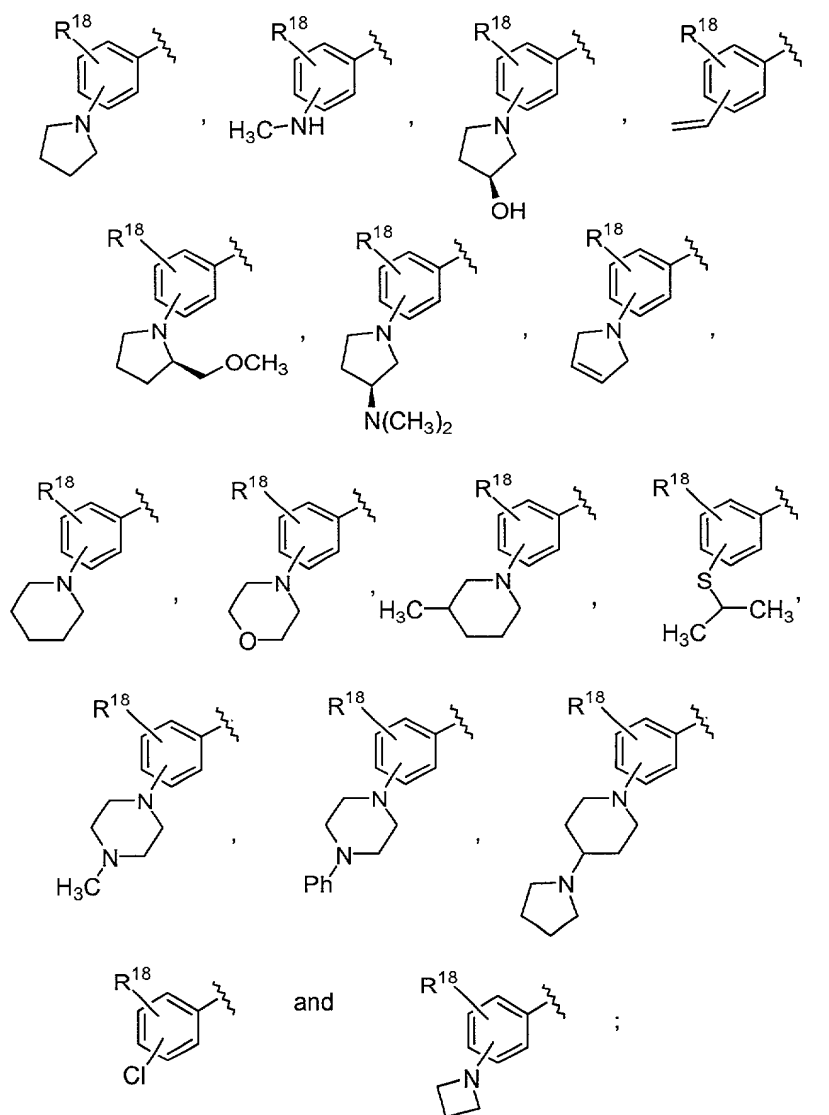
3

1 **30.** A composition in accordance with claim **29**, wherein A is a phenyl
2 group substituted with from one to three substituents selected from the group consisting
3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, nitro,
4 phenyl, naphthyl, pyrrolyl, pyrazolyl and -NR¹⁶R¹⁷ wherein R¹⁶ and R¹⁷ are
5 independently selected from the group consisting of hydrogen, (C₁-C₈)alkyl and (C₁-
6 C₈)heteroalkyl or are combined with the nitrogen atom to which each is attached to form
7 a four-, five-, six- or seven-membered ring optionally having additional heteroatoms as
8 ring members and optionally having additional substituents selected from the group
9 consisting of (C₁-C₈)alkyl, (C₁-C₈)heteroalkyl and phenyl.

1 **31.** A composition in accordance with claim **29**, wherein B is a phenyl
2 group substituted with from one to three substituents selected from the group consisting
3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy,
4 halogen, phenyl and phenoxy.

1 **32.** A composition in accordance with claim **29**, wherein A is a phenyl
2 group substituted with from one to three substituents selected from the group consisting
3 of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen and –
4 NR¹⁶R¹⁷ wherein R¹⁶ and R¹⁷ are independently selected from the group consisting of
5 hydrogen, (C₁-C₈)alkyl and (C₁-C₈)heteroalkyl or are combined with the nitrogen atom to
6 which each is attached to form a four-, five-, six- or seven-membered ring optionally
7 having additional heteroatoms as ring members and optionally having additional
8 substituents selected from the group consisting of (C₁-C₈)alkyl, (C₁-C₈)heteroalkyl and
9 phenyl, and B is a phenyl group substituted with from one to three substituents selected
10 from the group consisting of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-
11 C₄)haloalkyl, (C₁-C₄)haloalkoxy, halogen, phenyl and phenoxy.

1 **33.** A composition in accordance with claim **29**, wherein A is selected
2 from the group consisting of substituted or unsubstituted thienyl, substituted or
3 unsubstituted furanyl, substituted or unsubstituted indolyl, substituted or unsubstituted
4 benzothienyl, substituted or unsubstituted benzothienyl, and radicals of the formulae:



5
6
7 wherein
 R^{18} is a member selected from the group consisting of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)heteroalkyl, (C₁-C₄)haloalkyl, (C₁-C₄)haloalkoxy and halogen.

1 **34.** A composition in accordance with claim 33, wherein A is selected
2 from the group consisting of substituted or unsubstituted benzofuranyl, substituted or
3 unsubstituted benzothienyl, substituted or unsubstituted indolyl, substituted or
4 unsubstituted benzimidazolyl, substituted or unsubstituted benzthiazolyl and substituted
5 or unsubstituted benzoxazolyl.